

# United States Patent and Trademark Office



UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO. 9828
10/729,122	12/05/2003	James A. Williams	D-2939CIPCONDIV2	
33197	7590 10/07/2005		EXAM	INER
	KA, BUYAN & MULI	PORTNER, VIRGINIA ALLEN		
4 VENTURE IRVINE, CA	E, SUITE 300 N 92618		ART UNIT	PAPER NUMBER
, c. , 200			1645	

DATE MAILED: 10/07/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

# Notice of Non-Compliant Amendment (37 CFR 1.121)

Application No.	Applicant(s)		
10/729,122	WILLIAMS, JAMES A.		
Examiner	Art Unit		
Ginny Portner	1645		

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

The amendment document filed on  $\underline{6/13/05}$  is considered non-compliant because it has failed to meet the requirements of 37 CFR 1.121. In order for the amendment document to be compliant, correction of the following item(s) is required.

THE		DLLOWING MARKED (X) ITEM(S) CAUSE THE AMENDMENT DOCUMENT TO BE NON-COMPLIANT:  1. Amendments to the specification:  A. Amended paragraph(s) do not include markings.  B. New paragraph(s) should not be underlined.  C. Other
		2. Abstract:
		<ul><li>☐ A. Not presented on a separate sheet. 37 CFR 1.72.</li><li>☐ B. Other</li></ul>
		3. Amendments to the drawings:
		A. The drawings are not properly identified in the top margin as "Replacement Sheet," "New Sheet," or "Annotated Sheet" as required by 37 CFR 1.121(d).
		B. The practice of submitting proposed drawing correction has been eliminated. Replacement drawings showing amended figures, without markings, in compliance with 37 CFR 1.84 are required.
		C. Other
	$\boxtimes$	4. Amendments to the claims:
		A. A complete listing of all of the claims is not present.
		<ul> <li>B. The listing of claims does not include the text of all pending claims (including withdrawn claims)</li> <li>C. Each claim has not been provided with the proper status identifier, and as such, the individual status</li> </ul>
		of each claim cannot be identified. Note: the status of every claim must be indicated after its claim number by using one of the following status identifiers: (Original), (Currently amended), (Canceled),
		(Previously presented), (New), (Not entered), (Withdrawn) and (Withdrawn-currently amended).
		<ul> <li>□ D. The claims of this amendment paper have not been presented in ascending numerical order.</li> <li>□ E. Other: see attached narrative.</li> </ul>

For further explanation of the amendment format required by 37 CFR 1.121, see MPEP § 714 and the USPTO website at <a href="http://www.uspto.gov/web/offices/pac/dapp/opla/preognotice/officeflyer.pdf">http://www.uspto.gov/web/offices/pac/dapp/opla/preognotice/officeflyer.pdf</a>.

### TIME PERIODS FOR FILING A REPLY TO THIS NOTICE:

- 1. Applicant is given **no new time period** if the non-compliant amendment is an after-final amendment or an amendment filed after allowance. If applicant wishes to resubmit the non-compliant after-final amendment with corrections, the **entire corrected amendment** must be resubmitted within the time period set forth in the final Office action.
- 2. Applicant is given **one month**, or thirty (30) days, whichever is longer, from the mail date of this notice to supply the **corrected section** of the non-compliant amendment in compliance with 37 CFR 1.121, if the non-compliant amendment is one of the following: a preliminary amendment, a non-final amendment (including a submission for a request for continued examination (RCE) under 37 CFR 1.114), a supplemental amendment filed within a suspension period under 37 CFR 1.103(a) or (c), and an amendment filed in response to a *Quayle* action.

<u>Extensions of time</u> are available under 37 CFR 1.136(a) <u>only</u> if the non-compliant amendment is a non-final amendment or an amendment filed in response to a *Quayle* action.

### Failure to timely respond to this notice will result in:

**Abandonment** of the application if the non-compliant amendment is a non-final amendment or an amendment filed in response to a *Quayle* action; or

**Non-entry** of the amendment if the non-compliant amendment is a preliminary amendment or supplemental amendment.

Art Unit: 1645

## Response to Amendment

1. The reply filed on June 13, 2005 is not fully responsive to the prior Office Action because: All prior examined claims were drawn to SEQ ID NO 28, serotype A botulinum toxin and all of these claims have been canceled and all new claims have been submitted directed to serotype E botulinum toxin of SEQ Id Nos 50 and 52. In light of the fact that botulinum toxins are serologically distinct neurotoxins, and therefore independent and distinct species of inventions, Applicant has submitted claims directed to a different invention, as originally presented claims 25-38, were directed to serotype A, SEQ ID NO 28. Since the period for reply set forth in the prior Office action has expired, this application will become abandoned unless applicant corrects the deficiency and obtains an extension of time under 37 CFR 1.136(a).

The date on which the petition under 37 CFR 1.136(a) and the appropriate extension fee have been filed is the date for purposes of determining the period of extension and the corresponding amount of the fee. In no case may an applicant reply outside the SIX (6) MONTH statutory period or obtain an extension for more than FIVE (5) MONTHS beyond the date for reply set forth in an Office action. A fully responsive reply must be timely filed to avoid abandonment of this application.

### Election/Restrictions

2. Newly submitted claims 39-48 are directed to an invention that is independent or distinct from the invention originally claimed for the following reasons: All of the previously examined claims directed to SEQ Id NO 28, serotype A botulinum toxin have been canceled and all new claims directed to SEQ ID NO 50 and 52, which have not previously been examined, nor recited in any of the claims in the instant Application at the time of first action, SEQ ID Nos 50 and 52

being directed to serotype E botulinum toxin fusion proteins which is an independent and distinct species of invention not previously examined nor considered on the record (see Das Gupta et al (1972, reference attached herewith), which teaches botulinum toxin are serologically distinct neurotoxins and therefore define patentably independent and distinct subject matter

Page 3

Since applicant has received an action on the merits for the originally presented invention, this invention has been constructively elected by original presentation for prosecution on the merits. Accordingly, claims 39-48 are withdrawn from consideration as being directed to a non-elected invention. See 37 CFR 1.142(b) and MPEP § 821.03.

3. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Ginny Portner whose telephone number is (571) 272-0862. The examiner can normally be reached on M-F, alternate Fridays off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Lynette Smith can be reached on (571) 272-0864. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Vgp September 30, 2005